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INVESTOR IN PEOPLE

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Dated

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# Patents Form 1/77

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The  
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1 / 77

17 OCT 2002

## Request for grant of a patent

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17 OCT 2002

LONDON

The Patent Office

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1. Your reference **4-32731P1**
2. Patent application number **0224200.6**  
(The Patent Office will fill in this part) **02 E756677-7 D00524**  
**7700 0.00-0224200.6**
3. Full name, address and postcode of the or of each applicant  
(underline all surnames) **NOVARTIS AG**  
**LICHTSTRASSE 35**  
**4056 BASEL**  
**SWITZERLAND**  
  
Patent ADP number (if you know it)  
If the applicant is a corporate body, give the country/state of its incorporation **SWITZERLAND** **71254 87005**
4. Title of invention **Organic compounds**

Novartis Pharmaceuticals UK Ltd  
Patents and Trademarks  
Wimblehurst Road  
HORSHAM  
West Sussex  
RH12 5AB ADP No 0718522002

B.A. YORKE & CO  
CHARTERED PATENT AGENTS  
COOMB HOUSE, 7 ST. JOHN'S ROAD  
ISLEWORTH  
MIDDLESEX TW7 6NH

Patents ADP number (if you know it)

1800001

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Country	Priority application number (if you know it)	Date of filing (day/month/year)
7. If this application is divided or otherwise derived from an earlier UK application, give the number and the filing date of the earlier application
 

Number of earlier application	Date of filing (day/month/year)
8. Is a statement of inventorship and of right to grant of a patent required in support of this request? (Answer 'Yes' if:
  - a) any applicant named in part 3 is not an inventor, or
  - b) there is an inventor who is not named as an applicant, or
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(see note (d))

## Patents Form 1/77

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Continuation sheets of this form

Description 25

Claim(s) 2

Abstract 1

Drawing(s)

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Priority documents

Translations of priority documents

Statement of inventorship and right to grant of a patent (*Patents Form 7/77*)

Request for preliminary examination and search (*Patents Form 9/77*) One

Request for substantive examination (*Patents Form 10/77*)

Any other documents  
(please specify)

11.

I/We request the grant of a patent on the basis of this application

Signature

Date

B.A. Yorke & Co

B.A. Yorke & Co.

17 October 2002

12. Name and daytime telephone number of person to contact in the United Kingdom

Mrs. E. Cheetham  
020 8560 5847

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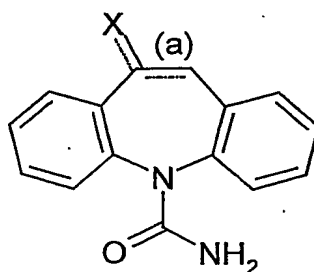
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ORGANIC COMPOUNDS

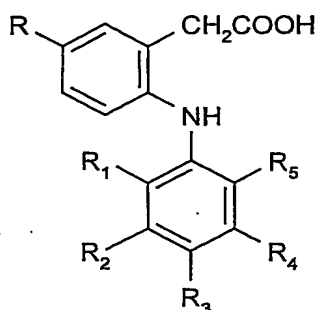
This invention relates to COX-2 inhibitors, in particular to combined use of COX-2 inhibitors with anti-epileptic compounds, and compositions containing such combinations.

which comprises in combination a carbamazepine or derivative thereof of formula I



I.

wherein X is =O, -OH or H, the bond between the azepine ring and X' being a double bond when X is =O and a single bond when X is -OH or H and the bond a of the azepine ring being a single bond when X is =O or -OH and a double bond when X' is H, and a COX-2 inhibitor of formula V



(V)

wherein R is methyl or ethyl;

R<sub>1</sub> is chloro or fluoro;

R<sub>2</sub> is hydrogen or fluoro;

R<sub>3</sub> is hydrogen, fluoro, chloro, methyl, ethyl, methoxy, ethoxy or hydroxy;

R<sub>4</sub> is hydrogen or fluoro; and

R<sub>5</sub> is chloro, fluoro, trifluoromethyl or methyl;

for simultaneous, sequential or separate use.

Further the invention provides the use of a COX-2 inhibitor of formula V as defined above for the preparation of a medicament, for use in combination with a carbamazepine or derivative thereof of formula II as defined above for treatment of pain.

In the alternative the invention provides use of a carbamazepine or derivative thereof of formula II as defined above for the preparation of a medicament for use in combination with a COX-2 inhibitor of formula V as defined above for treatment of pain.

In a further aspect the invention provides a method of treating a patient suffering from pain comprising administering to the patient an effective amount of a carbamazepine or derivative thereof of formula II as defined above and an effective amount of a COX-2 inhibitor of formula V as defined above.

In yet further aspects the invention provides:

- (i) A package comprising a carbamazepine or derivative thereof of formula II as defined above together with instructions for use in combination with a COX-2 inhibitor of formula V as defined above for treatment of pain, or
- (ii) A package comprising a COX-2 inhibitor of formula V as defined above together with instructions for use in combination with a carbamazepine or derivative thereof of formula II as defined above for treatment of pain.

Pain in general may be treated in accordance with the present invention including both nociceptive and inflammatory pain. In particular the combination treatment of the invention may be used for the treatment of musculoskeletal pain, especially lower back pain.

In the present description the terms "treatment" or "treat" refer to both prophylactic or preventative treatment as well as curative or disease modifying treatment, including treatment of patients at risk of suffering pain as well as patients who are already suffering pain.

Above and elsewhere in the present description the terms "an oxcarbazepine or derivative thereof" and "COX-2 inhibitor" include, as appropriate, pharmaceutically acceptable salts and esters thereof.

Particularly preferred compounds of formula V are those wherein R is methyl or ethyl; R<sub>1</sub> is chloro or fluoro; R<sub>2</sub> is hydrogen; R<sub>3</sub> is hydrogen, fluoro, chloro, methyl or hydroxy; R<sub>4</sub> is hydrogen; and R<sub>5</sub> is chloro, fluoro or methyl; pharmaceutically acceptable salts thereof; and pharmaceutically acceptable esters thereof.

A particularly preferred embodiment relates to the compounds of formula V wherein R is methyl or ethyl; R<sub>1</sub> is fluoro; R<sub>2</sub> is hydrogen; R<sub>3</sub> is hydrogen, fluoro or hydroxy; R<sub>4</sub> is hydrogen; and R<sub>5</sub> is chloro; pharmaceutically acceptable salts thereof; and pharmaceutically acceptable prodrug esters thereof.

Another particularly preferred embodiment of the invention relates to compounds of formula V wherein R is ethyl or methyl; R<sub>1</sub> is fluoro; R<sub>2</sub> is hydrogen or fluoro; R<sub>3</sub> is hydrogen, fluoro, ethoxy or hydroxy; R<sub>4</sub> is hydrogen or fluoro; and R<sub>5</sub> is chloro, fluoro or methyl; pharmaceutically acceptable salts thereof; and pharmaceutically acceptable prodrug esters thereof.

Further are said compounds wherein R is methyl or ethyl; R<sub>1</sub> is fluoro; R<sub>2</sub>-R<sub>4</sub> are hydrogen or fluoro; and R<sub>5</sub> is chloro or fluoro; pharmaceutically acceptable salts thereof; and pharmaceutically acceptable prodrug esters thereof.

A further embodiment of the invention relates to the compounds of formula V wherein R is methyl or ethyl; R<sub>1</sub> is fluoro; R<sub>2</sub> is fluoro; R<sub>3</sub> is hydrogen, ethoxy or hydroxy; R<sub>4</sub> is fluoro; and R<sub>5</sub> is fluoro; pharmaceutically acceptable salts thereof; and pharmaceutically acceptable prodrug esters thereof.

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Another embodiment of the invention relates to the compounds of formula V wherein R is methyl; R<sub>1</sub> is fluoro; R<sub>2</sub> is hydrogen; R<sub>3</sub> is hydrogen or fluoro; R<sub>4</sub> is hydrogen; and R<sub>5</sub> is chloro; pharmaceutically acceptable salts thereof; and pharmaceutically acceptable prodrug esters thereof.

Particularly preferred embodiments of the invention relate to compounds of formula V

(a) wherein R is methyl; R<sub>1</sub> is fluoro; R<sub>2</sub> is hydrogen; R<sub>3</sub> is hydrogen; R<sub>4</sub> is hydrogen; and R<sub>5</sub> is chloro; pharmaceutically acceptable salts thereof; and pharmaceutically acceptable prodrug esters thereof;

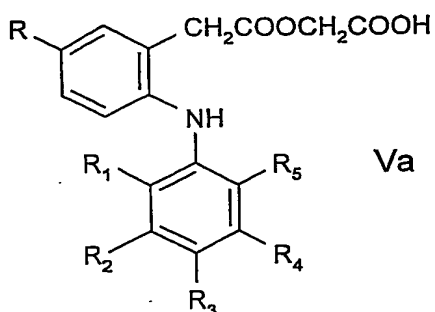
(b) wherein R is methyl; R<sub>1</sub> is fluoro; R<sub>2</sub> is hydrogen; R<sub>3</sub> is fluoro; R<sub>4</sub> is hydrogen; and R<sub>5</sub> is chloro; pharmaceutically acceptable salts thereof; and pharmaceutically acceptable prodrug esters thereof;

(c) wherein R is ethyl; R<sub>1</sub> is fluoro; R<sub>2</sub> is fluoro; R<sub>3</sub> is hydrogen; R<sub>4</sub> is fluoro; and R<sub>5</sub> is fluoro; pharmaceutically acceptable salts thereof; and pharmaceutically acceptable prodrug esters thereof; and

(d) wherein R is ethyl; R<sub>1</sub> is chloro; R<sub>2</sub> is hydrogen; R<sub>3</sub> is chloro; R<sub>4</sub> is hydrogen; and R<sub>5</sub> is methyl; pharmaceutically acceptable salts thereof; and pharmaceutically acceptable prodrug esters thereof.

Most preferably the COX-2 inhibitor of formula V is 5-methyl-2-(2'-chloro-6'-fluoroanilino)phenylacetic acid, or a salt or ester thereof.

Pharmaceutically acceptable prodrug esters of the compounds of formula V are ester derivatives which are convertible by solvolysis or under physiological conditions to the free carboxylic acids of formula V. Such esters are e.g. lower alkyl esters (such as the methyl or ethyl ester), carboxy-lower alkyl esters such as the carboxymethyl ester, nitrooxy-lower alkyl esters (such as the 4-nitrooxybutyl ester), and the like. Preferred prodrugs are the compounds of formula



wherein R and R<sub>1</sub>-R<sub>5</sub> have meaning as defined hereinabove for compounds of formula V; and pharmaceutically acceptable salts thereof.

Compounds of formula V and Va and their synthesis are described in published international patent applications Nos. WO 99/11605 and WO 01/23346, the teachings of which are incorporated herein by reference.

Carbamazepine is alternatively known as 5H-dibenz(b,f)azepine-5-carboxamide, G-32883, Biston, Calepsin, Carbelan, Epitol, Finlepsin, Sirtal, Stazepine, Tegretal, Tegretol, Telesmin and Timonil. The carbamazepine derivatives for use in the invention are alternatively known oxcarbazepine, GP-47680 and 10-oxo-10,11-dihydro-5H-dibenz(b,f)azepine-5-carboxamide (Trileptal<sup>®</sup>) and 10-hydroxy-10,11-dihydro-5H-dibenz(b,f)azepine-5-carboxamide, MHD or GP47779 respectively.

Monohydroxycarbamazepine (10-hydroxy-10,11-dihydro-carbamazepine), the main metabolite of the antiepileptic oxcarbazepine (Trileptal<sup>®</sup>) is well known from the literature [see for example Schuetz H. et al., Xenobiotica (GB), 16(8), 769-778 (1986)] and can be prepared synthetically starting from oxcarbazepine according to conventional methods.

Monohydroxycarbamazepine has been first disclosed in GB 1310120. The compound is indicated to be suitable for the treatment of psychosomatic disturbances, epilepsy, trigeminal neuralgia and cerebral spasticity.



Pharmacologically acceptable salts of carbamazepine, derivatives thereof and COX-2 inhibitors are preferably salts with bases, conveniently metal salts derived from groups Ia, Ib, IIa and IIb of the Periodic Table of the Elements, including alkali metal salts, e.g. potassium and especially sodium salts, or alkaline earth metal salts, preferably calcium or magnesium salts, and also ammonium salts with ammonia or organic amines.

The Agents of the Invention, i.e. the COX-2 inhibitor and the carbamazepine or derivative thereof are preferably used in the form of pharmaceutical preparations that contain the relevant therapeutically effective amount of of each active ingredient (either separately or in combination) optionally together with or in admixture with inorganic or organic, solid or liquid, pharmaceutically acceptable carriers which are suitable for administration. The Agents of the Invention may be present in the same pharmaceutical compositions, though are preferably in separate pharmaceutical compositions. Thus the active ingredients may be administered at the same time (e.g. simultaneously) or at different times (e.g. sequentially) and over different periods of time, which may be separate from one another or overlapping.

The pharmaceutical compositions may be, for example, compositions for enteral, such as oral, rectal, aerosol inhalation or nasal administration, compositions for parenteral, such as intravenous or subcutaneous administration, or compositions for transdermal administration (e.g. passive or iontophoretic).

The particular mode of administration and the dosage may be selected by the attending physician taking into account the particulars of the patient, especially age, weight, life style, activity level, and disease state as appropriate

Preferably, both the COX-2 inhibitor and carbamazepine or derivative pharmaceutical compositions are adapted for oral or parenteral (especially oral) administration. Intravenous and oral, first and foremost oral, administration is considered to be of particular importance. Preferably the COX-2 inhibitor active ingredient is in oral form.

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The dosage of COX-2 inhibitor administered is dependent on the species of warm-blooded animal (mammal), the body weight, age and individual condition, and on the form of administration. A unit dosage for oral administration to a mammal of about 50 to 70 kg may contain between about 5 and 1500 mg, e.g. from 100-1000 mg, preferably 200-800 mg of the active ingredient.

COX-2 inhibitor formulations in single dose unit form contain preferably from about 1% to about 90%, and formulations not in single dose unit form contain preferably from about 0.1% to about 20%, of the active ingredient. Single dose unit forms such as capsules, tablets or dragées contain e.g. from about 1mg to about 1500mg of the active ingredient.

COX-2 inhibitor formulations in single dose unit form contain preferably from about 1% to about 90%, and formulations not in single dose unit form contain preferably from about 0.1% to about 20%, of the active ingredient. Single dose unit forms such as capsules, tablets or dragées contain e.g. from about 1mg to about 1500mg of the active ingredient.

Similarly the dosage of carbamazepine or derivative administered is dependent on the species of warm-blooded animal (mammal), the body weight, age and individual condition, and on the form of administration. In general, the daily dosage of carbamazepine or derivative varies between about 3 mg/kg and about 20 mg/kg. Suitable unit dosage forms, such as dragées, tablets or suppositories, preferably contain 30-200mg of carbamazepine or derivative. Dosage units for oral administration preferably contain between 10% and 90% by weight of carbamazepine or derivative.

Pharmaceutical preparations for enteral and parenteral administration are, for example, those in dosage unit forms, such as dragées, tablets or capsules and also ampoules. They are prepared in a manner known *per se*, for example by means of conventional mixing, granulating, confectioning, dissolving or lyophilising processes. For example, pharmaceutical preparations for oral administration can be obtained by combining the active ingredient with solid carriers, where

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appropriate granulating a resulting mixture, and processing the mixture or granulate, if desired or necessary after the addition of suitable adjuncts, into tablets or dragée cores.

Other orally administrable pharmaceutical preparations are dry-filled capsules made of gelatin, and also soft, sealed capsules made of gelatin and a plasticiser, such as glycerol or sorbitol. The dry-filled capsules may contain the active ingredient in the form of a granulate, for example in admixture with fillers, such as lactose, binders, such as starches, and/or glidants, such as talc or magnesium stearate, and, where appropriate, stabilisers. In soft capsules the active ingredient is preferably dissolved or suspended in suitable liquids, such as fatty oils, paraffin oil or liquid polyethylene glycols, it being possible also for stabilisers to be added.

Parenteral formulations are especially injectable fluids that are effective in various manners, such as intravenously, intramuscularly, intraperitoneally, intranasally, intradermally or subcutaneously. Such fluids are preferably isotonic aqueous solutions or suspensions which can be prepared before use, for example from lyophilised preparations which contain the active ingredient alone or together with a pharmaceutically acceptable carrier. The pharmaceutical preparations may be sterilised and/or contain adjuncts, for example preservatives, stabilisers, wetting agents and/or emulsifiers, solubilisers, salts for regulating the osmotic pressure and/or buffers.

Suitable formulations for transdermal application include an effective amount of the active ingredient with carrier. Advantageous carriers include absorbable pharmacologically acceptable solvents to assist passage through the skin of the host. Characteristically, transdermal devices are in the form of a bandage comprising a backing member, a reservoir containing the compound optionally with carriers, optionally a rate controlling barrier to deliver the active ingredient of the skin of the host at a controlled and predetermined rate over a prolonged period of time, and means to secure the device to the skin.

The following examples are intended to illustrate the invention and are not to be construed as being limitations thereon.

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EXAMPLES

## A. Formulation Examples

Example 1

Table 1

Ingredient	Amount per 200 mg tablet batch (kg)
<b>Core</b>	
<b>Granulation</b>	
5-methyl-2-(2'-chloro-6'-fluoroanilino)phenylacetic acid drug substance	50**
Microcrystalline cellulose, NF (PH 101)	12.85
Lactose monohydrate, NF	11.65
Croscarmellose sodium, NF	1
Povidone, USP	4
Titanium dioxide, USP	2
Water, purified ***, USP	20.375
<b>Extra-granular Phase</b>	
Microcrystalline cellulose, NF (PH 102)	13
Croscarmellose sodium, NF	3
Titanium dioxide, USP	2
Magnesium stearate, NF	0.5
<b>Coating</b>	
Opadry white	2.801 ****
Opadry yellow	2.0 ****
Opadry red	0.4 ****
Opadry black	0.0504 ****
Water, purified ***, USP	29.758 ****

\*\* The weight of drug substance is taken with reference to the dried substance (100-per-cent) on the basis of the assay value (factorization). The difference in weight is adjusted by the amount of microcrystalline cellulose used.

\*\*\* Removed during processing.

\*\*\*\* Includes a 50 % excess for loss during the coating process.

Table 1, above, sets out the formula for a batch of approximately 250,000 immediate release film-coated tablets of 5-methyl-2-(2'-chloro-6'-fluoroanilino)-phenylacetic acid. To make the tablets, titanium dioxide is dispersed in water, followed by the addition of povidone and mixing for 20 minutes to make a povidone/titanium dioxide suspension. The drug substance, lactose, microcrystalline cellulose, and croscarmellose are mixed in a high shear mixer (e.g., a Collette Gral) for 5 minutes to form a drug mixture. The drug mixture is granulated in the high shear mixer with the povidone/titanium dioxide suspension. The suspension is pumped at a rate of 3 kg/min into the drug mixture. The resulting mixture is mixed an additional 90 seconds after all the suspension is added. The wet granulation is dried in a fluid bed dryer, using an inlet air temperature of 50 °C. The residual water target is 3.5 % (with a permissible range of 2.5 – 4.5 %). The dried granulation is passed through a screen using a mill (oscillator) and a 30 mesh screen. The previous steps are repeated to make a second granulation.

The extra-granular phase titanium dioxide is passed through a 60 mesh hand screen. The dry granulations are mixed with the extra-granular phase microcrystalline cellulose, croscarmellose sodium and titanium dioxide in a twin shell mixer for 300 revolutions to form a penultimate mixture. Magnesium stearate is passed through a 60 mesh hand screen and is mixed with the penultimate mixture in a twin shell mixer for 50 revolutions to form a tableting mixture. The tableting mixture is pressed into tablets using a tablet press and oval punches.

The coating powders (Opadry) are mixed with purified water to make a 15 % w/w coating suspension. The tablets are film coated with the coating suspension in a coating pan using 60 °C to 75 °C inlet air temperature.

Table 2 sets out the contents of a 200 mg 5-methyl-2-(2'-chloro-6'-fluoroanilino)phenylacetic acid film-coated tablet.

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Table 2

Ingredient	Theoretical amount [mg]	Function
<b>Core</b>		
5-methyl-2-(2'-chloro-6'-fluoroanilino)phenylacetic acid drug substance	200	Active substance
Microcrystalline cellulose (PH 101)	51.4	Filler
Lactose	46.6	Filler
Povidone	16	Binder
Titanium dioxide	8	Color
Croscarmellose sodium	4	Disintegrant
Water, purified *	Q.S.	Granulating liquid
<b>Extragranular phase</b>		
Microcrystalline cellulose (PH 102)	52	Filler
Croscarmellose sodium	12	Disintegrant
Titanium dioxide	8	Color
Magnesium stearate	2	Lubricant
Core weight	400	
<b>Coating</b>		
Opadry white (00F18296)	7.4676	Color
Opadry yellow (00F12951)	5.3312	Color
Opadry red (00F15613)	1.0668	Color
Opadry black (00F17713)	0.1344	Color

Ingredient	Theoretical amount [mg]	Function
Water, purified *	Q.S.	Coating solvent
<b>Total weight</b>	<b>414</b>	

\* removed during processing

In addition, the tablet formulations may contain 5-methyl-2-(2'-chloro-6'-fluoroanilino)benzyl alcohol and/or 5-methyl-2-(2'-chloro-6'-fluoroanilino)benzoic acid in an amount between about 0.01 and 2% by weight, more specifically between about 0.1 and 1

Example 2

An alternative formulation is as set out in Table 3, with information about as percentage w/w, mg/dose, and kg/ 50,000 tablet batch.

(a) Table 3 Alternative formulation composition

% w/w	Ingredient	Mg/dose	Kg/batch
Granulation			
65.04	5-methyl-2-(2'-chloro-6'-fluoroanilino) phenylacetic acid drug substance	400.00	20.00
2.15	Croscarmellose sodium, NF (Ac-Di-Sol)	13.22	0.661
6.60	Povidone K30, USP	40.59	2.029
18.12	Purified water, USP*	Qs	Qs
Blending			
23.56	Microcrystalline Cellulose, NF (Avicel PH 102)	144.90	6.066
2.15	Croscarmellose sodium, NF (Ac-Di-Sol)	13.22	0.553
0.50	Magnesium Stearate, NF (vegetable source)	3.07	0.128
Film Coating			
84.46	Opadry, Global White 00F18296	15.2028	0.296637
14.03	Opadry, Global Red 00F15613	2.5254	0.049275
1.51	Opadry, Global Black 00F17713	0.2718	0.005303
	Purified Water, USP*	Qs	1.990218
Film Coated Tablet Weight		633.00	

\*Does not appear in final product. Percentage of water added used for granulation based on the dry weight of drug substance and croscarmellose sodium.

The batch is granulated as described in Example 1. The granulation is dried to residual moisture (% LOD) of 1.79%. The formulation process is the same as for the development batches as described above, except for the additional step of coating with Opadry in a coating pan. The coating powders (Opadry) are mixed with purified water to make a 15 % w/w coating suspension.

The tablets are film coated with the coating suspension in a coating pan using 60°C to 75°C inlet air temperature. Based on friability data, a target force of 18 KN (16 – 20 KN range) is used to



compress the remainder of the batch, resulting in acceptable friability (less than 0.5%) and the disintegration times of less than 5 mins. The ejection force is approximately 800 N throughout the compression run. This demonstrates that the blend is lubricated adequately. No picking/sticking is observed on the punch surfaces after 225 minutes. Thus, a smaller size tablet with high drug loading (65%) is achieved using a high shear granulation process, using 17 X 6.7 mm ovaloid tooling to get tablets with acceptable hardness and friability characteristics.

In addition, the tablet formulations may contain 5-methyl-2-(2'-chloro-6'-fluoroanilino)benzyl alcohol and/or 5-methyl-2-(2'-chloro-6'-fluoroanilino)benzoic acid in an amount between about 0.01 and 2% by weight, more specifically between about 0.1 and 1%.

### Example 3

#### Wet granulated tablet composition

<u>Amount per tablet</u>	<u>Ingredient</u>
25 mg	COX-2 inhibitor
79.7 mg	Microcrystalline cellulose
79.7 mg	Lactose monohydrate
6 mg	Hydroxypropyl cellulose
8 mg	Croscarmellose sodium
0.6 mg	Iron oxide
1 mg	Magnesium stearate

Tablet dose strengths of between 5 and 125 mg can be accommodated by varying total weight, and the ratio of the first three ingredients. Generally it is preferable to maintain a 1:1 ratio for microcrystalline cellulose: lactose monohydrate.

### Example 4

#### Hard gelatine capsule composition

<u>Amount per capsule</u>	<u>Ingredient</u>
25 mg	COX-2 inhibitor

37 mg Microcrystalline cellulose  
 37 mg Lactose anhydrate  
 1 mg Magnesium stearate  
 1 capsule Hard gelatin capsule

Capsule dose strengths of between 1 and 50 mg can be accommodated by varying total fill weight, and the ratio of the first three ingredients. Generally it is preferable to maintain a 1:1 ratio for microcrystalline cellulose:lactose monohydrate.

#### Example 5

Oral solution

#### Amount per 5mL Ingredient

50 mg COX-2 inhibitor  
 to 5 mL with Polyethylene oxide 400

#### Example 6

#### Intravenous infusion

#### Amount per 200 mL dose Ingredient

1 mg COX-2 inhibitor  
 0.2 mg Polyethylene oxide 400  
 1.8 mg Sodium chloride  
 to 200 mL Purified water

#### Example 7:

Oxcarbazepine Formulations

Tablet-core	[mg]	[mg]
TRILEPTAL AS (oxcarbazepine)	300.0	600.0

extra fine		
Cellulose HPM 603	8.4	16.8
Microcrystalline Cellulose	65.6	131.2
Colloidal anhydrous silica	1.6	3.2
Magnesium stearate	4.4	8.8
Crosspovidone	20.0	40.0
<b>Core Weight</b>	<b>400.0</b>	<b>800.0</b>

<b>Coating</b>		
Cellulose HPM 603	7.351	11.946
Iron(II) oxide (yellow) 17268	0.499	0.811
Polyethylene glycol (PEG) 8000	1.331	2.162
Talcum	5.323	8.649
Titanium dioxide	1.497	2.432
<b>Coating Weight</b>	<b>16.000</b>	<b>26.000</b>

Oxcarbazepine, cellulose HPM 603 (binder) and a portion (approximately half) of the microcrystalline cellulose (binder, filler, disintegration-promoting excipient) are mixed in a mixer, preferably in a high-speed mixer, e.g. DIOSNA, LOEDIGE, FIELDER or GLATT. Water is added to the mixture and the mixture kneaded, preferably in a high-speed mixer until an adequate consistency is achieved. Alternatively, the HPM 603 may be dissolved in the water, beforehand. The mixture is formed into wet granules using an ALEXANDER Reibschnitzler, QUADRO-COMILL and the granules dried in a fluidised bed (AEROMATIC, GLATT). The remainder of the microcrystalline cellulose is added together with AEROSIL 200 (flow conditioner) and crospovidone (disintegrator) to the dry granules followed by mixing in a comminuter (FREWITT, QUADRO-COMILL, FITZMILL). Finally, magnesium stearate (lubricant) is added with mixing (STOECKLIN container mixer, VRIECO mixer). Alternatively, the lubricant may be added directly to the comminuted material. The final mixture is compressed to form TRILEPTAL tablets (eccentric press, rotary press: KILIAN, KORSCH, FETTE, MANESTY).

The tablets are coated with an aqueous preparation consisting of cellulose HPM 603 (film former), iron oxide yellow 17268 (pigment), PEG 8000 (plasticiser for the film former), talcum (anti-adhesive agent, covering agent) and titanium dioxide (covering agent) in a rotating coating pan (ACCELA-COTA, GLATT, DRIACOATER, DUMOULIN). Alternatively, it is possible to use, for example, fluidised-bed or air-suspension apparatus for the coating process (AEROMATIC, GLATT, FREUND, HUETTLIN).

### Example 8

An alternative formulation is as set out in Table 5, with information about as percentage w/w, mg/dose, and kg/ 50,000 tablet batch.

<b>Example 1</b>	(mg)	(mg)	(mg)
<b>Tablet Core:</b>			
Oxcarbazepine	150	300	600
Avicel PH 102	32,8	65,6	131,2
Cellulose HPM 603	4,2	8,4	16,8
Polyvinylpyrrolidone	10	20	40
Aerosil 200	0,8	1,6	3,2
Magnesium stearate	2,2	4,4	8,8
	<b>200</b>	<b>400</b>	<b>800</b>
<b>Coating:</b>			
Polyethylene glycol (PEG) 8000	0,832	1,331	2,162
Cellulose HPM 603	4,595	7,352	11,947
Talcum	3,327	5,323	8,649

Titanium Dioxide	0,935	1,496	2,431
Iron oxide, yellow	0,312	0,499	0,81
	10	16	26
Total	210	416	826

Mix the TRILEPTAL, cellulose HPM 603 (binder) and AVICEL PH 102 (binder, filler, disintegration-promoting excipient) in a mixer, preferably in a high-speed mixer (DIOSNA, LOEDIGE, FIELDER, GLATT etc.). Add water as granulation liquid to the mixture, and knead in a mixer, preferably a high-speed mixer, until an adequate consistency is achieved. Alternatively, the binder cellulose HPM may be dissolved in the granulation liquid, water, beforehand. Granulate the wet granules using a suitable device (ALEXANDER Reibschneider, QUADRO-COMILL) and dry in a fluidised bed (AEROMATIC, GLATT). Add AVICEL PH 102, AEROSIL 200 (flow conditioner) and polyvinylpyrrolidone PXL (disintegrator) to the dry granules and comminute and mix in a comminuter (FREWITT, QUADRO-COMILL, FITZMILL). Finally, add magnesium stearate (lubricant) and mix (STOECKLIN container mixer, VRIECO mixer). Alternatively, the lubricant may be added directly to the comminuted material. Compress the final mixture to form TRILEPTAL tablets (eccentric press, rotary press: KILIAN, KORSCH, FETTE, MANESTY).

Coat the tablets with an aqueous preparation consisting of cellulose HPM 603 (film former), iron oxide yellow 17268 (pigment), PEG 8000 (plasticiser for the film former), talcum (anti-adhesive agent, covering agent) and titanium dioxide (covering agent) in a rotating coating pan (ACCELA-COTA, GLATT, DRIACOATER, DUMOULIN). Alternatively, it is possible to use, for example, fluidised-bed or air-suspension apparatus for the coating process (AEROMATIC, GLATT, FREUND, HUETTLIN).

#### Example 9:

A typical oral solution of oxcarbazepine comprises (% expressed in mass/volume):

Oxcarbazepine, micronized, fine)	1 to 20%	(e.g. TRILEPTAL/AS, extra
Avicel RC 591	0.1 to 1.9%	
Methylparaben	0.01 to 1%	
Polyethylene glycol 400 monostearate	0.01 to 1%	
Propylene glycol (1,2-propanediol), dist.	0.5 to 10%	
Propylparaben	0.005 to 0.5%	
Saccharin sodium, cryst.	0.005 to 0.5 %	
Sorbic acid	0.005 to 0.5 %	
Sorbitol solution (non crystallizing)	10 to 40%	
Ascorbic acid	0.1 to 10%	
Water, purified	40 to 85%	
Yellow plum-lemon aroma	0 to 15%	

**Example 10:**

Parenteral Formulation with oxcarbazepine or a derivative thereof of formula I: Oxcarbazepine or a derivative thereof is dissolved under a nitrogen blanket with stirring at 60- 80° C. in water for injection (WFI) at a concentration of 2.5 mg/ml. After cooling to room temperature anhydrous glucose for injection is added and dissolved by stirring under nitrogen purging to obtain a 4.75% concentration glucose in water. After filtration through 0.22 micrometer pore size filter, the solution is purged with nitrogen, filled in glass vials (class II quality), sealed with a rubber closure and alu-cap and sterilized by autoclaving at 121° C. for 15 minutes. The vials are stable and clear of coloured particles for at least two years at 2-8° C.

**Example 11:**

Alternative parenteral Formulation with oxcarbazepine or a derivative thereof of formula I: Oxcarbazepine or a derivative thereof dissolved with stirring at 60-80° C. in WFI at a concentration of 2.5 mg/ml. After cooling to room temperature glucose for injection (anhydrous) is added and dissolved by stirring to obtain a 4.75% concentration of glucose in water. After filtration through 0.22 micrometer pore size filter, the solution is filled in glass vials, sealed with a rubber closure and alu-cap and sterilized by autoclaving at 121° C. for 15 minutes.

The vials are stable and clear of coloured particles for at least three months at 2-8° C.

**Example 12:**

Alternative parenteral Formulation with oxcarbazepine or a derivative thereof of formula I:  
Oxcarbazepine or a derivative thereof is dissolved under a nitrogen blanket by stirring at 60- 80° C. in WFI at a concentration of 2.5 mg/ml. After cooling to room temperature sodium chloride is added and dissolved with stirring under nitrogen purging to obtain a 0.9% concentration of sodium chloride in water. After filtration through 0.22 micrometer pore size filter, the solution is purged with nitrogen, filled in glass vials, sealed with a rubber closure and alu-cap and sterilized by autoclaving at 121° C. for 15 minutes.

The vials are inspected after three months storage at 2-8° C. and show the presence of red coloured particles.

**Example 13:**

Alternative parenteral Formulation with oxcarbazepine or a derivative thereof of formula I:  
Oxcarbazepine or a derivative thereof is dissolved with stirring at 60-80° C. in WFI at a concentration of 2.5 mg/ml. After cooling to room temperature sodium chloride is added and dissolved by stirring to obtain a 0.9% concentration of sodium chloride in water. After filtration through 0.22 micrometer pore size filter, the solution is filled in glass vials, sealed with a rubber closure and alu-cap and sterilized by autoclaving at 121° C. for 15 minutes. The solutions in the vials show within six weeks storage at 2-8° C. the presence of red coloured particles.

**Example 14:**

Carbamazepine formulations:

**Core**

Carbamazepine – water free (Tegretol®)	200	mg
Microcrystalline Cellulose (Avicel® – FMC – Corporation Philadelphia)	20	mg
Hydroxypropylmethyl cellulose (Pharmacoat® 603 – Shin-Etsu Chem. Co. – Tokyo)	12,5	mg
Vinylpyrrolidone/Vinylacetate-60:40-Copolymers (Kollidon® VA 64 – BASF Ludwigshafen)	80	mg

Polyethylenglycol (MG:5 x 10 <sup>6</sup> -Polyox® - Coagulant – Union Carbide)	80	mg
Sodium chloride (purest)	80	mg
Sodium laurylsulfate	6	mg
Magnesium stearate	11,5	mg
	=	490 mg

**Coating**

Cellulose acetate (32,0)	16	mg
Cellulose acetate (39,8)	20	mg
Polyethylenglycol 4000	4	mg
	=	40 mg
<hr/> Total weight		530 mg

**Alternative Formulation:****Core**

Carbamazepine – water free (Tegretol®)	200	mg
Microcrystalline cellulose (Avicel® – FMC – Corporation Philadelphia)	20	mg
Hydroxypropylmethyl cellulose (Pharmacoat® 603 – Shin-Etsu Chem. Co. – Tokyo)	13	mg
Vinylpyrrolidone/Vinylacetate-60:40-Copolymers (Kollidon® VA 64 – BASF Ludwigshafen)	80	mg
Hydroxyethyl cellulose (Tylose H 4000 PHA)	80	mg
Glucose (purest)	90	mg
Sodium laurylsulfate	7	mg
Magnesium stearate	10	mg
	=	500 mg

**Coating**

<del>Cellulose acetate (32,0)</del>	<del>16</del>	<del>mg</del>
Cellulose acetate (39,8)	20	mg



Polyäthylenglycol 4000	4	mg	
		=	40 mg
Total weight			540 mg

**Alternative Formulation:****Core**

Carbamazepine – water free (Tegretol®)	200	mg	
Microcrystalline cellulose (Avicel® – FMC – Corporation Philadelphia)	20	mg	
Hydroxypropylmethylcellulose (Pharmacoat® 603 – Shin-Etsu Chem. Co. – Tokyo)	12,5	mg	
Vinylpyrrolidon/Vinylacetate-60:40-Copolymers (Kollidon® VA 64 – BASF Ludwigshafen)	81,3	mg	
Polyethylenglycol (MG:5 x 10 <sup>6</sup> -Polyox® - Coagulant – Union Carbide)	80	mg	
Sodium lauryl sulfate	6	mg	
Magnesium stearate	10,2	mg	
		=	410 mg

**Coating**

Cellulose acetate (32,0)	16	mg	
Cellulose acetate (39,8)	20	mg	
Polyethylenglycol 4000	4	mg	
		=	40 mg

Total weight	450 mg
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**Alternative Formulation:****Core**

Carbamazepine – water free (Tegretol®)	200	mg
Hydroxypropylmethyl cellulose (Pharmacoat® 603 –		

Shin-Etsu Chem. Co. – Tokyo)	25	mg	
Hydroxyethyl cellulose (Natrosol® - 250L – Hercules)	25	mg	
Hydroxyethyl cellulose (Natrosol® - 250H)	25	mg	
Mannit	215	mg	
Sodium lauryl sulfate	5	mg	
Magnesium stearat	5	mg	
	=		500 mg

**Coating**

Cellulose acetate (32,0)	18,9	mg	
Cellulose acetate (39,8)	2,9	mg	
Hydroxypropylmethyl cellulose 15 CPS	2,1	mg	
Polyethylenglycol 8000	2,1	mg	
	=		26 mg
<hr/> Total weight			526 mg

**Example 14**     **Treatment of Patients****Assumptions:**

- 1) Two formulations :     200 mg Prexige plus 300 mg Trileptal  
    200 mg Prexige plus 600 mg Trileptal
- 2) b.i.d. dosing
- 3) limited titration
- 4) effective dose Trileptal = 900 - 1200 mg/day  
     effective dose Prexige = 400 mg/day
- 5) sample size would have to be estimated by a statistician
- 6) trial timeline is set up to achieve POC but possibly not statistical significance

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Design: double-blind, placebo-controlled, parallel group, multicenter

Duration: 4 to 6 weeks including screening

Patient population:

*inclusion criteria* - male or female  $\geq$  to 18 years old

- low back pain (below T6 and above gluteal fold) that may radiate to leg
- pain lasting more than three months
- pain present on five out of seven days
- VAS score  $\geq$  to 40 mm on four of the last seven days
- comprehensive history and physical examination including focused neurological examination

*exclusion criteria* - unstable spinal segment

- progressive neurological deficits
- excluded drugs: all other NSAIDs, opioids, TCAs, AEDs, oral steroids except for treatment of asthma or skin conditions, steroid injections
- other pain conditions that may interfere with assessment of the low back pain
- patients previously treated with either Prexige or Trileptal
- patients with hypersensitivity to carbamazepine, oxcarbazepine or lumiracoxib and other non-steroidal anti-inflammatories including aspirin
- patients with active disability compensation claims or any litigation related to their radiculopathic pain.

Variables:

*primary efficacy variable* - VAS

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*secondary efficacy variables* - responder rate, sleep assessment, SF-36, POMS, assessment of back mobility and low-back pain specific QOL

## Suggested visit schedule:

visit 1 (day -14 to day - 1)	screening
visit 2 (day 1)	randomization, titration and treatment
visit 3 (day 21)	withdrawal
visit 4 (day 28)	final visit

The withdrawal phase can be eliminated to give 4 weeks total treatment (1 week titration, 3 weeks maintenance).

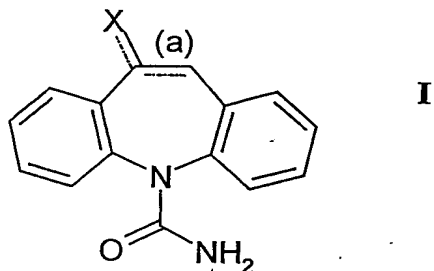
## Titration and maintenance dosing schedule:

Day	AM Dose <sup>a</sup>	PM Dose <sup>a</sup>	Total Daily Doses <sup>a</sup>
1	0	200/300	200/300
2	200/300	200/300	400/600
3	200/300	200/300	400/600
4	200/300	200/600	400/900
5	200/300	200/600	400/900
6	200/600	200/600	400/1200
7-21	200/600	200/600	400/1200
22-28	0	0	0

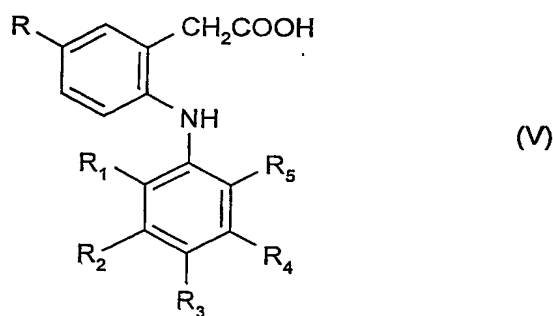
<sup>a</sup>expressed as mg Prexige/mg Trileptal

CLAIMS

1. A pharmaceutical composition for treatment of pain, which comprises in combination oxcarbazepine or derivative thereof of formula I



wherein X is =O, -OH or H, the bond between the azepine ring and X' being a double bond when X is =O and a single bond when X is -OH or H and the bond a of the azepine ring being a single bond when X is =O or -OH and a double bond when X' is H, and a COX-2 inhibitor of formula V



wherein R is methyl or ethyl;

R<sub>1</sub> is chloro or fluoro;

R<sub>2</sub> is hydrogen or fluoro;

R<sub>3</sub> is hydrogen, fluoro, chloro, methyl, ethyl, methoxy, ethoxy or hydroxy;

R<sub>4</sub> is hydrogen or fluoro; and

R<sub>5</sub> is chloro, fluoro, trifluoromethyl or methyl;

for simultaneous, sequential or separate use.

2. Use of a COX-2 inhibitor of formula V as defined in claim 1 for the preparation of a medicament, for use in combination with oxcarbazepine or derivative thereof of formula I as defined in claim 1, for treatment of pain.
  3. Use of oxcarbazepine or derivative thereof of formula I as defined in claim 1, for the preparation of a medicament for use in combination with a COX-2 inhibitor of formula V as defined in claim 1 for treatment of pain.
  4. A method of treating a patient suffering from pain comprising administering to the patient an effective amount of oxcarbazepine or derivative thereof of formula I as defined above, and an effective amount of a COX-2 inhibitor of formula V as defined in claim 1.
  5. A package comprising oxcarbazepine or derivative thereof of formula I as defined in claim 1, together with instructions for use in combination with a COX-2 inhibitor of formula V as defined in claim 1 for treatment of pain, or a package comprising a COX-2 inhibitor of formula V as defined in claim 1 together with instructions for use in combination with oxcarbazepine or derivative thereof of formula I as defined in claim 1, for treatment of pain.
  6. A composition method, use or package according to any one of the preceding claims in which the COX-2 inhibitor is 5-methyl-2-(2'-chloro-6'-fluoroanilino)phenylacetic acid, or a pharmaceutically acceptable salt or ester thereof.
  7. A composition method, use or package according to any one of claims 1 to 6 in which the carbamazepine derivative is oxcarbazepine.
  8. A composition method, use or package according to any one of claims 1 to 6 in which the ~~carbamazepine derivative is 10-hydroxy-10,11-dihydro-5H-dibenz(b,f)azepine-5-~~carboxamide.
-

ABSTRACT

ORGANIC COMPOUNDS

A pharmaceutical composition for treatment of pain, comprises in combination carbamazepine or derivative thereof as defined and a COX-2 inhibitor as defined for simultaneous, sequential or separate use. Also provided is a method of treating a patient suffering from pain, comprising administering to the patient an effective amount of carbazmaepine or derivative thereof as defined and an effective amount of a COX-2 inhibitor as defined.

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